Dkt. #706-A-US



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants : JIANG, Shibo and DEBNATH, Asim Kumar

U.S. Serial No.: 10/706,027

Filed Date : November 12, 2003

For : COMPOUNDS FOR INHIBITION OF HIV INFECTION

BY BLOCKING HIV ENTRY

Law Offices of Albert Wai-Kit Chan, LLC

World Plaza, Suite 604 141-07 20th Avenue Whitestone, NY 11357

December 16, 2003

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir/Madam:

INFORMATION DISCLOSURE STATEMENT

In accordance with their duty of disclosure under 37 C.F.R. §1.56, Applicants would like to direct the Examiner's attention to the following references which are listed below and on PTO/SB/08B (Exhibit A), with each individual reference further attached as Exhibit 1 through 14.

- Chan , D. C., C. T. Chutkowski , and P. S. Kim . 1998. Evidence that a prominent cavity in the coiled coil of HIV type 1 gp41 is an attractive drug target. Proc. Natl . Acad. Sci . U S A 95:15613-15617. [Exhibit 1]
- Chan , D. C., D. Fass , J. M. Berger , and P. S. Kim . 1997. Core structure of gp41 from the HIV envelope glycoprotein. Cell 89:263-273. [Exhibit 2]
- 3. Debnath, A. K., L. Radigan, and S. Jiang. 1999. Structure-based identification of small molecule antiviral compounds targeted to the gp41 core

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structure of the human immunodecifiency virus type 1. J. Med. Chem. 42:3203-3209. [Exhibit 3]

- 4. Eckert, D. M., V. N. Malashkevich, L. H. Hong, P. A. Carr, and P. S. Kim . 1999. Inhibiting HIV-1 entry: discovery of D-peptide inhibitors that target the gp41 coiled-coil pocket. Cell 99:103-115. [Exhibit 4]
- 5. Ernst, J. T., O. Kutzki, A. K. Debnath, S. Jiang, H. Lu, and A. D. Hamilton. 2002. Design of a Protein Surface Antagonist Based on alpha-Helix Mimicry: Inhibition of gp41 Assembly and Viral. Angew. Chem. Int. Ed Engl. 41:278-281. [Exhibit 5]
- 6. Jiang, S. and A. K. Debnath. 2000. A salt bridge between an N-terminal coiled coil of gp41 and an antiviral agent targeted to the gp41 core is important for anti-HIV-1 activity. Biochem. Biophys. Res. Commun. 270:153-157. [Exhibit 6]
- 7. Jiang, S., K. Lin, and M. Lu. 1998. A conformation-specific monoclonal antibody reacting with fusion-active gp41 from the HIV-1 envelope glycoprotein. J. Virol. 72:10213-10217. [Exhibit 7]
- 8. Jiang, S., K. Lin, N. Strick, and A. R. Neurath. 1993. HIV-1 inhibition by a peptide. Nature 365:113. [Exhibit 8]
- 9. Jiang, S., K. Lin, L. Zhang, and A. K. Debnath. 1999. A screening assay for antiviral compounds targeted to the HIV-1 gp41 core structure using a conformationspecific monoclonal antibody. J. Virol. Methods 80:85-96. [Exhibit 9]
- 10. Jiang, S., Q. Zhao, and A. K. Debnath. 2002. Peptide
 and Non-peptide HIV Fusion Inhibitors. Curr. Pharm.
 Des. 8:563-580. [Exhibit 10]
- 11. Lin, P. F., W. Blair, T. Wang, T. Spicer, Q. Guo, N. Zhou, Y. F. Gong, H. G. Wang, R. Rose, G. Yamanaka, B. Robinson, C. B. Li, R. Fridell, C. Deminie, G. Demers, Z. Yang, L. Zadjura, N. Meanwell, and R. Colonno.

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2003. A small molecule HIV-1 inhibitor that targets the HIV-1 envelope and inhibits CD4 receptor binding. Proc. Natl. Acad. Sci. U. S. A 100:11013-11018. [Exhibit 11]

- 12. Liu, S., Q. Zhao, and S. Jiang. 2003. Determination of the HIV-1 gp41 postfusion conformation modeled by synthetic peptides: applicable for identification of the HIV-1 fusion inhibitors. Peptide In press. [Exhibit 12]
- 13. Weissenhorn , W., A. Dessen , S. C. Harrison , J. J. Skehel , and D. C. Wiley . 1997. Atomic Structure of the Ectodomain from HIV-1 gp41. Nature 387:426-428. [Exhibit 13]
- 14. Zhao, Q., J. T. Ernst, A. D. Hamilton, A. K. Debnath, and S. Jiang. 2002. XTT formazan widely used to detect cell viability inhibits HIV type 1 infection in vitro by targeting gp41. AIDS Res. Hum. Retroviruses 18:989-997. [Exhibit 14]

If a telephone interview would be of assistance in advancing prosecution of the subject application, Applicants' undersigned attorney invites the Examiner to telephone him at the number provided below.

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No fee is deemed necessary in connection with the filing of this Information Disclosure Statement. However, if any additional fee is required, authorization is hereby given to charge the amount of any such fee to Deposit Account No. 50-1891.

I hereby certify that this paper is being deposited this date with the U.S. Postal Service with sufficient postage for first class mail addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450

Albert Wai-Kit Chan Date
Reg. No. 36,479

Respectfully submitted,

Albert Wai-Kit Chan Registration No. 36,479 Attorney for Applicants Law Offices of Albert Wai-Kit Chan, LLC World Plaza, Suite 604 141-07 20th Avenue Whitestone, New York 11357

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PTO/SB/08B (04-03) Approved for use through 04/30/2003. OMB 0651-0031

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Numb r	10/706,027	
			CLOSURE	Filing Dat	November 12, 2003	
			PPLICANT	First Nam d Inv ntor	JIANG, Shibo	
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Sheet	1	of	2	Attorney Docket Number	706-A-US	
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		NON PATENT LITERATURE DOCUMENTS	
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
	1	Chan , D. C., C. T. Chutkowski , and P. S. Kim . 1998. Evidence that a prominent cavity in the coiled coil of HIV type 1 gp41 is an attractive drug target. Proc. Natl . Acad. Sci . U S A 95:15613-15617.	
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	4	Eckert, D. M., V. N. Malashkevich, L. H. Hong, P. A. Carr, and P. S. Kim . 1999. Inhibiting HIV-1 entry: discovery of D-peptide inhibitors that target the gp41 coiled-coil pocket. Cell 99:103-115.	
	5	Ernst, J. T., O. Kutzki, A. K. Debnath, S. Jiang, H. Lu, and A. D. Hamilton. 2002. Design of a Protein Surface Antagonist Based on alpha-Helix Mimicry: Inhibition of gp41 Assembly and Viral Fusion. Angew. Chem. Int. Ed Engl. 41:278-281.	
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	10	Jiang, S., Q. Zhao, and A. K. Debnath. 2002. Peptide and Non-peptide HIV Fusion Inhibitors. Curr. Pharm. Des. 8:563-580.	

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This collection of information is required by 37 CFR 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 120 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete his form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents. Washington, DC 20231. Commissioner for Patents, Washington, DC 20231.

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			PPLICANT	First Named Inventor	JIANG, Shibo	
<i>"</i>				Art Unit		
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Sheet	2	of	2	Attorney Docket Number	706-A-US	

		NON PATENT LITERATURE DOCUMENTS	
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	11	Lin, P. F., W. Blair, T. Wang, T. Spicer, Q. Guo, N. Zhou, Y. F. Gong, H. G. Wang, R. Rose, G. Yamanaka, B. Robinson, C. B. Li, R. Fridell, C. Deminie, G. Demers, Z. Yang, L. Zadjura, N.	
		Meanwell, and R. Colonno. 2003. A small molecule HIV-1 inhibitor that targets the HIV-1 envelope and inhibits CD4 receptor binding. Proc. Natl. Acad. Sci. U. S. A 100:11013-11018.	
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